

The role of surgical cytoreduction in the treatment of malignant pleural mesothelioma: Meeting summary of the International Mesothelioma Interest Group Congress, September 11-14, 2012, Boston, Mass

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The treatment of all solid tumors, including malignant pleural mesothelioma (MPM), is dependent on (1) macroscopic complete resection and (2) treatment of micrometastatic disease. The role of surgery in the treatment of MPM has been the subject of debate after the recent publication of the Mesothelioma and Radical Surgery (MARS) I trial.¹ The International Mesothelioma Interest Group (IMIG) met from September 11 through 14, 2012, in Boston, Mass. During this meeting, more than 500 participants representing all the involved specialty groups met in multiple comprehensive sessions to review, critique, and extend the state of knowledge regarding the role of surgery, including both extended pleurectomy/decortication (P/D) and extrapleural pneumonectomy (EPP), in the treatment of MPM.

Some of the deficiencies of the MARS I trial, which was published a year ago in *Lancet Oncology*, were discussed in multiple sessions of the IMIG meeting. The editorial that accompanied the publication articulated numerous shortcomings of the trial.² The MARS I trial was designed as a pilot feasibility trial, the result of which was negative in that it failed to demonstrate the feasibility of randomly allocating patients to surgery versus no surgery. Nevertheless, the

publication contained an analysis of tertiary end points, including survival, which was based on the small pilot cohort, representing fewer than 10% of the required sample size for an adequately powered between-arm comparison as published by the MARS trialists. Protocol compliance was also poor in that 6 of 26 patients in the no EPP group underwent off-protocol surgery, whereas only 16 of 24 patients in the EPP group actually underwent EPP.

Quality control of the surgery in the MARS trial, if undertaken, was not reported. Intent-to-treat morbidity (11/24; 46%) and mortality (3/24; 13%), and more strikingly, EPP-associated morbidity (11/16; 69%) and mortality (3/16; 19%), were much higher than reported in the literature. The chemotherapy regimens applied were uncontrolled. Neither final histologic type nor disease stage was reported for the patients who underwent surgery, leaving an open question as to whether these patients, who demonstrated survival inferior to most previous reports, may have had disproportionate N2 or nonepithelial disease. Conversely, the reported 19-month median survival among chemotherapy-only (no EPP) patients was clearly anomalous when compared with a vast prospective literature. The long-term outcome of the study cohort remains unknown, because the overall survival analysis was truncated at 18 months, whereas the quality of life data were reported to 24 months. These deficiencies make drawing any conclusions from MARS I regarding the therapeutic efficacy of EPP impossible.

The patterns of failure in MPM were reaffirmed at the 2012 IMIG meeting. Dr Elizabeth H. Baldini, in reference to her previous work, presented a contemporary group of patients and demonstrated essentially the same distribution of recurrence as originally reported, which is primarily local.³ Six institutional series from the US, Europe, and Japan involving macroscopic complete resection by EPP or P/D in the setting of multimodality treatment of MPM were presented at the meeting.⁴⁻⁹ These reports were discussed in detail in light of previous literature to date. Median survival ranged from 25 to 37 months for patients with epithelial disease and negative extrapleural lymph nodes. Operative mortality ranged from 0% to 2%.

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On behalf of the International Association for the Study of Lung Cancer (IASLC), Dr Valerie Rusch presented a preliminary analysis of the IASLC staging project, which has since been published in the November 2012 issue of the *Journal of Thoracic Oncology*.¹⁰ In the IASLC worldwide registry of patients with all stages of epithelial MPM, the analysis showed 19-month median survival among 1359 patients undergoing surgical resection (P/D or EPP). Moreover, patients undergoing EPP for early-stage disease demonstrated survival superior to that of all other subgroups, a median of 40 months. On the basis of the current literature and the IASLC report, it was concluded by IMIG members that surgery, whether P/D or EPP, with the goal of obtaining a macroscopic complete resection should be performed in the multimodality treatment of MPM. In particular, it was agreed that the type of cytoreductive procedure should be selected on the basis of disease distribution, institutional experience, and surgeon preference and experience. Furthermore, it was collectively decided that these operations should be performed by surgeons who have achieved morbidity and mortality within the scope of the current literature.

After much discussion in multiple forums and settings with surgeons, medical oncologists, radiation oncologists, epidemiologists, and basic scientists, the attendees of the 2012 IMIG meeting reached agreement on the following points:

- Surgical macroscopic complete resection and control of micrometastatic disease play a vital role in the multimodality therapy of MPM, as is the case for other solid malignancies.
- Surgical cytoreduction is indicated when macroscopic complete resection is deemed achievable.
- The type of surgery (EPP or P/D) depends on clinical factors and on individual surgical judgment and expertise.
- All patients with the diagnosis of MPM should be initially evaluated in a multidisciplinary setting, including medical oncology, radiation oncology, and surgery.
- Clinical staging (lymph node sampling, positron emission tomography, magnetic resonance imaging) should be performed before therapy.
- The histologic subtype should be identified by tissue biopsy before initiation of therapy.

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